## **CLAIMS AMENDMENT**

1. (currently amended): A compound of the formula

$$A_{n}^{1}A_{n}^{2}A_{n}^{3}A^{4}A^{5}A^{6}A^{7}A^{8}A^{9}A^{10}A^{11}A^{12}A^{13}A^{14}A^{15}A^{16}A^{17}A^{18}$$

$$A_{n}^{1}A_{n}^{2}A_{n}^{3}A^{4}AA_{n}^{6}ICA_{n}^{9}QIA_{n}^{12}YA_{n}^{14}FGA_{n}^{17}F$$

$$(1)$$

and acylated and/or amidated forms thereof,

wherein each n is independently 0 or 1;

A<sup>1</sup>, A<sup>2</sup>[[,]] and A<sup>3</sup> are each independently any amino acid;

A<sup>4</sup>, A<sup>12</sup>[[,]] and A<sup>17</sup> are independently acidic amino acids E, D or Q;

[[A<sup>13</sup>,]] A<sup>14</sup>[[, A<sup>15</sup>, and A<sup>18</sup> are independently]] is an aromatic amino [[acids]] acid;

A<sup>5</sup>, A<sup>7</sup>, A<sup>8</sup>, A<sup>11</sup>, and A<sup>16</sup> represent any amino acid;

 $A^{6}[[,]]$  and  $A^{9}[[, and A^{10}]]$  represent independently a basic amino acid or a polar neutral amino acid;

wherein each of said amino acids may be in the L form, racemic form, or D form, with the proviso that

the compound of formula (1) is other than does not comprise

AALEAKICHQIEYYFGDF ALEAKICHQIEYYFGDF when all amino acids are in the

L-form, and

must be in isolated form when all amino acids are in the L-form and formula (1) is of the sequence ALEAKICHQIEYYFGDF, AALEAKICHQIEYYFGDF, LDLDTKICEQIEYYFGDF, AALEAKICHQIEEYYFGDF, DDADQRIIKQLEYYFGNI, VSKLEASTIRQEYYFGDA or QERAIIRQVEYYFGDF.

- 2. (original): The compound of claim 1 wherein all amino acids are gene encoded.
- 3. (currently amended): The compound of claim 1 wherein all linkages between A<sup>i</sup> subunits the amino acids are amide linkages.
- 4. (currently amended): The compound of claim 1 where <u>in</u> all of [[A<sup>i</sup>]] <u>the amino</u> acids are in the D form.

- 5. (currently amended): The compound of claim 1 wherein all of [[A<sup>i</sup>]] the amino acids are in the L form.
- 6. (original): The compound of claim 1 wherein each of A<sup>4</sup>, A<sup>12</sup> and A<sup>17</sup> is independently aspartic or glutamic.
- 7. (currently amended): The compound of claim 1 wherein each of A<sup>13</sup>, A<sup>14</sup>[[, A<sup>15</sup> and A<sup>18</sup>]] is independently phenylalanine or tyrosine.
  - 8. (canceled)
- 9. (currently amended): The compound of claim 1 wherein each of A<sup>6</sup>[[,]] and A<sup>9</sup>[[and A<sup>10</sup>]] is independently lysine, histidine, arginine, glutamine, or asparagine.
- 10. (previously presented): The compound of claim 1 which is selected from the group consisting of AALEAQICQQIEYYFGDF (SEQ ID NO:2), AALQAKICHQIQYYFGQF (SEQ ID NO:3), QQQEAKICHQIEYYFGDF (SEQ ID NO:4) and AALEAKICHQIEYQFGDF (SEQ ID NO:12).
- 11. (currently amended): The compound of claim 1 which is in isolated or purified form and is selected from the group consisting of ALEAKICHQIEYYFGDF (SEQ ID NO:13), AALEAKICHQIEYYFGDF (SEQ ID NO:14), LDLDTKICEQIEYYFGDF (SEQ ID NO:15), AALEAKICHQIEEYYFGDF (SEQ ID NO:16), DDADQRIIKQLEYYFGNI (SEQ ID NO:17), VSKLEASTIRQEYYFGDA (SEQ ID NO:18) and QERAIIRQVEYYFGDF (SEQ ID NO:19).
- 12. (original): A pharmaceutical, veterinary or agricultural/horticultural composition which comprises the compound of claim 1 along with a suitable excipient.

13-19. (canceled)

- 20. (withdrawn): A method to treat viral infection in a plant or animal subject which method comprises administering to said subject an antivirally effective amount of the compound of claim 1.
- 21. (withdrawn): The method of claim 20 wherein said method further comprises administering at least one additional antiviral agent.
- 22. (withdrawn): The method of claim 21 wherein said administering of the compound and said at least one additional antiviral agent is substantially simultaneous.
- 23. (withdrawn): The method of claim 21 wherein said administering of the compound of claim 1 and said at least one antiviral compound is sequential.
- 24. (withdrawn): The method of claim 21 wherein said additional antiviral compound is I-RNA.

25-35. (canceled)